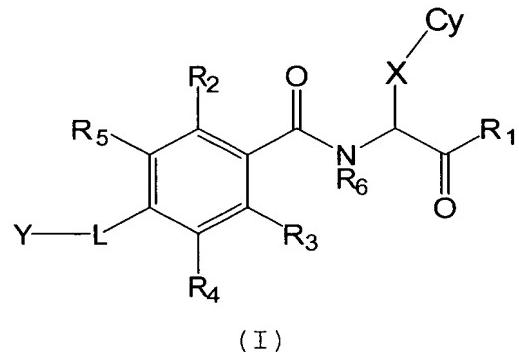


1. A compound of formula (I)



wherein

10 Cy is a non-aromatic carbocycle or heterocycle optionally substituted with hydroxyl, mercapto, thioalkyl, halogen, oxo, thio, amino, aminoalkyl, amidine, guanidine, nitro, alkyl, alkoxy or acyl;

15 X is a divalent hydrocarbon chain optionally substituted with hydroxyl, mercapto, halogen, amino, aminoalkyl, nitro, oxo or thio and optionally interrupted with N, O, S, SO or SO₂;

20 Y is a carbocycle or heterocycle optionally substituted with hydroxyl, mercapto, halogen, oxo, thio, thioalkyl, amino, aminoalkyl, carbocycle or heterocycle ring, hydrocarbon, a halo-substituted hydrocarbon, amino, amidine, guanidine, cyano, nitro, alkoxy or acyl;

25 L is a bond or a divalent hydrocarbon chain optionally substituted hydroxyl, halogen, oxo or thio and optionally interrupted with N, O, S, SO or SO₂ or an amino acid residue; less than 3 or 5 atoms

30 R₁ is H, OH, amino, O-carbocycle or alkoxy optionally substituted with amino, a carbocycle or heterocycle;

5 R₂₋₅ are independently H, hydroxyl, mercapto, halogen, cyano, amino, amidine, guanidine, nitro or alkoxy; or R₃ and R₄ together form a fused carbocycle or heterocycle optionally substituted with hydroxyl, halogen, oxo, thio, amino, amidine, guanidine or alkoxy;

10 R₆ is H or a hydrocarbon chain optionally substituted with a carbocycle or a heterocycle; and salts, solvates and hydrates thereof;

15 with the proviso that when Y is phenyl, R₂, R₄ and R₅ are H, R₃ is Cl and R₁ is OH then X is other than cyclohexyl.

2. A compound according to claim 1, wherein Cy is a 5- or 6-member non-aromatic heterocycle optionally substituted with hydroxyl, mercapto, thioalkyl

20 halogen, oxo, thio, amino, aminoalkyl, amidine, guanidine, nitro, alkyl, alkoxy or acyl.

3. A compound according to claim 2, wherein said

25 heterocycle comprises one or two heteroatoms and is optionally substituted with hydroxyl, oxo, mercapto, thio, alkyl or alkanoyl.

4. A compound according to claim 3, wherein said

30 heterocycle is selected from the group consisting of piperidine, piperazine, morpholine, tetrahydrofuran, tetrahydrothiophene, oxazolidine, cyclopropapyrrolidine and thiazolidine optionally substituted with hydroxy, oxo, mercapto, thio, alkyl or alkanoyl.

35 5. A compound according to claim 4, wherein said heterocycle is selected from the group consisting of

- 5 piperidine, piperazine, morpholine, tetrahydrofuran,
tetrahydrothiophene, oxazolidine, thiazolidine
optionally substituted with hydroxy, oxo, mercapto,
thio, alkyl or alkanoyl.
- 10 6. A compound according to claim 1, wherein Cy is a 3-6
member carbocycle optionally substituted with
hydroxyl, mercapto, halogen, oxo, thio, amino,
amidine, guanidine, alkyl, alkoxy or acyl.
- 15 7. A compound according to claim 6, wherein said
carbocycle is partially unsaturated.
8. A compound according to claim 7, wherein Cy is
cyclopropyl, cyclypropenyl, cyclobutyl, cyclbutenyl,
20 cyclopentyl, cyclopentenyl cyclohexyl or
cyclohexenyl.
9. A compound according to claim 1, wherein X is a C₁₋₅
divalent hydrocarbon optionally having one or more
25 carbon atoms replaced with N, O, S, SO or SO₂ and
optionally being substituted with hydroxyl, oxo or
.thio.
10. A compound according to claim 1, wherein X is -CH₂-
30 NR₆-C(O)- wherein the carbonyl -C(O)- portion thereof
is covalently bound to Cy and R₆ is H or alkyl.
11. A compound according to claim 1, wherein Y is a
carbocycle or heterocycle optionally substituted
35 with hydroxyl or halogen.
12. A compound according to claim 11, wherein Y is
furan-2-yl, thiophene-2-yl or phenyl, wherein said

5 phenyl is optionally substituted with halogen or hydroxyl.

- 10
13. A compound according to claim 1, wherein L is a divalent hydrocarbon optionally having one or more carbon atoms replaced with N, O, S, SO or SO₂ and optionally being substituted with hydroxyl, halogen oxo or thio; or three carbon atoms of the hydrocarbon are replaced with an amino acid residue.
- 15 14. A compound according to claim 13, wherein L is -CH=CH-C(O)-NR₆-CH₂-, -CH₂-NR₆-C(O)-, -C(O)-N₆-CH₂-, -CH(OH)-(CH₂)₂-, -(CH₂)₂-CH(OH)-, -(CH₂)₃-, -C(O)-NR₆-CH(R₇)-C(O)-NR₆-, -NR₆-C(O)-CH(R₇)-NR₆-C(O)-, -CH(OH)-CH₂-O- or -CH(OH)-CF₂-CH₂- wherein each R₆ is independently H or alkyl and R₇ is an amino acid side chain.
- 20
15. A compound according to claim 14, wherein R₁ is H, OH, amino, O-carbocycle or alkoxy optionally substituted with a carbocycle.
- 25
16. A compound according to claim 15, wherein R₁ is H or C₁₋₄ alkyloxy.
- 30 17. A compound according to claim 1, wherein at least one of R₂ and R₃ is halogen and the other is H or halogen.
- 35
18. A compound according to claim 17, wherein R₂ and R₃ are both Cl.
19. A compound according to claim 18, wherein R₄ and R₅ are both H.

20. A pharmaceutical composition comprising a compound according to claim 1 with a pharmaceutically acceptable adjuvant, diluent or carrier.
- 10 21. A method of inhibiting binding of a LFA-1 to a protein ligand comprising contacting LFA-1 with a compound of claim 1.
22. A method of treating a disease or condition mediated by LFA-1 in a mammal comprising administering to said mammal an effective amount of a compound according to claim 1.
23. A method according to claim 23, wherein said disease or condition is arthritis, psoriasis, organ transplant rejection, asthma, and inflammatory bowel disease
- 15 23. A method of inhibiting an inflammatory disease or condition in a mammal comprising administering to said mammal an effective amount of a compound according to claim 1.